

Our Ref.: AB-392 (F2001-129)

TITLE OF THE INVENTION

CURABLE COMPOSITION, CURED COATING FILM AND COATED
SUBSTRATE

5

BACKGROUND

FIELD OF THE INVENTION

The present invention relates to a curable
composition capable of forming a cured coating film
excellent in abrasion resistance, transparency, anti-
10 fouling property and non-tackiness (release property), a
cured coating film formed by such a curable composition
and a coated substrate having such a cured coating film.

DISCUSSION OF BACKGROUND

A protective film to be formed on the surface of a
15 substrate such as a lens, a display or a photomask, is
required have a hard coat property, an anti-fouling
property, non-tackiness, etc.

However, heretofore, it used to be required to carry
out a hard coat processing to form a hard coating film
20 and an anti-fouling non-tackiness processing to form a
coating film having a low surface tension property such
as water repellency or oil repellency, separately, in
order to form a protective film having the above-
mentioned plurality of properties on the surface of a
25 substrate. Accordingly, separate steps are required,
thus leading to an increase of costs.

In order to reduce the costs, various attempts have

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been proposed to form a coating film satisfying the above-mentioned plurality of properties, in a single step. For example, JP-A-10-104403 discloses a sheet having a cured film made of an ultraviolet curable resin and an oligomer containing a fluorine atom-containing alkylalkoxysilane; JP-A-6-211945 discloses a sheet having a cured film made of an alkyl fluoroacrylate and an acrylate not-compatible with an alkyl fluoroacrylate; and JP-A-11-305420 discloses a protective layer for a photomask having a release property from a resist improved.

However, in the case of JP-A-10-104403, the reactivity between the ultraviolet curable resin and the fluorine atom-containing oligomer is poor, whereby there was a problem that the fluorine atom-containing oligomer tends to bleed out as time passes. In the case of JP-A-6-211945, the compatibility of the two materials is poor, whereby there was a problem that a uniform coating film can not be obtained, and it is difficult to obtain a coating film having transparency and good appearance. And, in the case of JP-A-11-305420, a heat curing step is required, whereby there was a problem that the productivity is low.

SUMMARY OF THE INVENTION

It is an object of the present invention to provide a curable composition capable of forming, in a single coating step, a cured coating film which has an anti-

fouling property and a release property and is excellent in abrasion resistance, whereby such properties will not be deteriorated even when used for a long period of time, a cured coating film formed by such a curable

5 composition, and a coated substrate having such a cured coating film.

The present invention provides a curable composition comprising a compound (A) having at least one active energy ray curable polymerizable functional group, a
10 fluorine-containing copolymer (B) obtained by (1) copolymerizing a polymerizable monomer (a) having a polyfluoroalkyl group and a polymerizable monomer (b) having a photo-curable functional group, or (2) introducing a photo-curable functional group into a
15 fluorine-containing copolymer (D) obtained by copolymerizing a polymerizable monomer (a) having a polyfluoroalkyl group and a polymerizable monomer (d) having a group capable of introducing a photo-curable functional group, and a photopolymerization initiator
20 (C).

In the present invention, the curable composition preferably contains from 0.01 to 20 parts by mass of the fluorine-containing copolymer (B) and from 0.01 to 20 parts by mass of the photopolymerization initiator (C),
25 per 100 parts by mass of the compound (A).

Further, the present invention provides a cured coating film formed by the above curable composition.

Still further, the present invention provides a coated substrate comprising a substrate and such a cured coating film formed on at least one side of the substrate. In this invention, the substrate is preferably a photomask.

According to the present invention, it is possible to provide a curable composition capable of forming, in a single coating step, a cured coating film which has an anti-fouling property and a release property and is excellent in abrasion resistance, whereby such properties will not deteriorate even when used for a long period of time, a cured coating film having the above-mentioned properties, formed by such a curable composition, and a coated substrate having such a cured coating film.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Now, the components constituting the curable composition of the present invention will be described. In the following description, an acryloyl group and an methacryloyl group will be generally referred to as a (meth)acryloyl group. The same applies to such a representation as a (meth)acryloyloxy group, a (meth)acrylic acid or a (meth)acrylate.

Firstly, the compound (A) having at least one active energy ray curable polymerizable functional group (hereinafter referred to simply as the compound (A)) will be described.

Among compounds (A), a compound having one

polymerizable functional group which is polymerizable by
an active energy ray (hereinafter referred to simply as a
monofunctional compound) is preferably a compound having
a (meth)acryloyl group, particularly preferably a
5 compound having an acryloyl group. It may further have
other functional groups such as a hydroxyl group and an
epoxy group. The following compounds may, for example,
be mentioned as such a monofunctional compound.

An alkyl (meth)acrylate (the carbon number of the
10 alkyl group: 1 to 13), allyl (meth)acrylate, benzyl
(meth)acrylate, butoxyethyl (meth)acrylate, butanediol
(meth)acrylate, butoxytriethyleneglycol
mono(meth)acrylate, t-butylaminoethyl (meth)acrylate, 3-
chloro-2-hydroxypropyl (meth)acrylate, 2-cyanoethyl
15 (meth)acrylate, cyclohexyl (meth)acrylate,
dicyclopentanyl (meth)acrylate, 2,3-dibromopropyl
(meth)acrylate, dicyclopentenyl (meth)acrylate,
dicyclopentenylloxyethyl (meth)acrylate, N,N-
diethylaminoethyl (meth)acrylate, N,N-dimethylaminoethyl
20 (meth)acrylate, 2-ethoxyethyl (meth)acrylate, 2(2-
ethoxyethoxy)ethyl (meth)acrylate, 2-ethylhexyl
(meth)acrylate, glycerol (meth)acrylate, glycidyl
(meth)acrylate, heptadecafluorodecyl (meth)acrylate, 2-
hydroxyethyl (meth)acrylate, 2-hydroxy-3-
25 (meth)acryloyloxypropyltrimethylammonium chloride, 2-
hydroxypropyl (meth)acrylate, isobonyl (meth)acrylate, 3-
(meth)acryloyloxypropyltrimethoxysilane, 2-methoxyethyl

(meth)acrylate, methoxydiethyleneglycol (meth)acrylate,
methoxytriethyleneglycol (meth)acrylate,
methoxytetraethyleneglycol (meth)acrylate,
methoxydipropylene glycol (meth)acrylate, methoxylated
5 cyclodecatriene (meth)acrylate, morpholine
(meth)acrylate, nonylphenoxypolyethylene glycol
(meth)acrylate, nonylphenoxypolypropylene glycol
(meth)acrylate, octafluoropentyl (meth)acrylate,
phenoxyhydroxypropyl (meth)acrylate, phenoxyethyl
10 (meth)acrylate, phenoxydiethylene glycol (meth)acrylate,
phenoxytetraethylene glycol (meth)acrylate,
phenoxyhexaethylene glycol (meth)acrylate, phenoxy
(meth)acrylate, polypropylene glycol (meth)acrylate,
sodium sulfonate ethoxy (meth)acrylate, tetrafluoropropyl
15 (meth)acrylate, tetrahydrofurfuryl (meth)acrylate,
trifluoroethyl (meth)acrylate, vinyl acetate, N-vinyl
caprolactam, N-vinyl pyrrolidone, N-vinyl formamide,
morpholino (meth)acrylate, and 2-morpholinoethyl
(meth)acrylate.

20 Further, as a compound having two or more
polymerizable functional groups curable by an active
energy ray (hereinafter referred to simply as a
polyfunctional compound), compounds disclosed in
paragraphs 0016 to 0020, and 0023 to 0047, in JP-A-11-
25 240103, may preferably be mentioned.

As a preferred polyfunctional compound, a compound
having two or more (preferably from 2 to 50, more

preferably from 3 to 30) polymerizable functional groups of at least one type selected from (meth)acryloyl groups, may be mentioned. Among them, a compound having two or more (meth)acryloyloxy groups, i.e. a polyester of a compound having two or more hydroxyl groups such as a polyhydric alcohol, with (meth)acrylic acid, is preferred. Further, it may be a compound having various functional groups or bonds in addition to the above-mentioned polymerizable functional group. Particularly preferred are a (meth)acryloyl group-containing compound having a urethane bond (hereinafter referred to as an acrylurethane) and a (meth)acrylate compound having no urethane bond.

The above acrylurethane may, for example, be 1) a polyfunctional compound which is an acrylurethane as a reaction product of pentaerythritol or a polypentaerythritol as its oligomer, a polyisocyanate and a hydroxyalkyl (meth)acrylate, and which has at least three (more preferably from 4 to 20) polymerizable functional groups curable by an active energy ray, or 2) a polyfunctional compound which is an acrylurethane as a reaction product of a hydroxyl group-containing poly(meth)acrylate of pentaerythritol or polypentaerythritol, with a polyisocyanate, and which has at least three (more preferably from 4 to 20) polymerizable functional groups curable by an active energy ray.

As the (meth)acrylate compound having no urethane bond, a pentaerythritol type poly(meth)acrylate or an isocyanurate type poly(meth)acrylate may be mentioned. Here, the pentaerythritol type poly(meth)acrylate means a polyester of pentaerythritol or polypentaerythritol with (meth)acrylic acid (which preferably has from 4 to 20 polymerizable functional groups curable by an active energy ray). The isocyanurate type poly(meth)acrylate means a polyester of a tris(hydroxyalkyl) isocyanurate or a compound obtained by adding from 1 to 6 mol of caprolactam or an alkylene oxide to 1 mol of a tris(hydroxyalkyl) isocyanurate, with (meth)acrylic acid (which preferably has 2 or 3 polymerizable functional groups curable by an active energy ray).

In the present invention, the above-mentioned preferred polyfunctional compound may be used in combination with another polyfunctional compound having at least two polymerizable functional groups curable by an active energy ray (particularly a poly(meth)acrylate of a polyhydric alcohol).

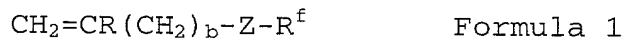
In the compound (A), the proportion of the above-mentioned monofunctional compound or the above-mentioned polyfunctional compound, is not particularly limited.

Now, the fluorine-containing copolymer (B) will be described.

The fluorine-containing copolymer (B) in the present invention, is a compound obtained by (1) copolymerizing a

polymerizable monomer (a) having a polyfluoroalkyl group
(hereinafter referred to simply as a polymerizable
monomer (a)) and a polymerizable monomer (b) having a
photo-curable functional group (hereinafter referred to
5 simply as a polymerizable monomer (b)), or (2)
introducing a photo-curable functional group into a
fluorine-containing copolymer (d) obtained by
copolymerizing the polymerizable monomer (a) and a
polymerizable monomer (d) having a group capable of
10 introducing a photo-curable functional group (hereinafter
referred to simply as a polymerizable monomer (d)).

As the polymerizable monomer (a), a compound of the
following formula 1 is preferred.



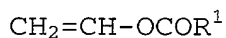
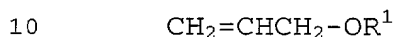
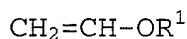
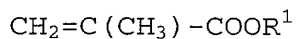
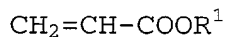
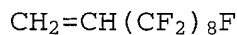
15 wherein R is a hydrogen atom or a C₁₋₄ alkyl group, b is
an integer of from 0 to 6, Z is a single bond or a group
selected from the group consisting of -CH₂-, -O-, -COO-,
-CONH-, -NHCO- and -OCO-, and R^f is a C₂₋₄₀
polyfluoroalkyl group or a C₂₋₄₀ polyfluoroalkenyl group,
20 provided that at least one hydrogen atom in such a group
may be substituted by a hydroxyl group or a halogen atom,
and an etheric oxygen atom or a thioetheric sulfur atom
may be inserted between a carbon-carbon bond.

R is preferably a hydrogen atom or a methyl group.

25 R^f is preferably a C₂₋₂₂ polyfluoroalkyl group, more
preferably a C₄₋₁₈ polyfluoroalkyl group, most preferably
a C₆₋₁₄ polyfluoroalkyl group. Particularly preferred is

a C₆₋₁₄ linear perfluoroalkyl group wherein all hydrogen atoms in the alkyl group are substituted by fluorine atoms.

Specifically, the following compounds may be mentioned as preferred as the polymerizable monomer (a):



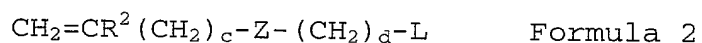
In the above formulae, R¹ represents the following groups (m: an even number of from 4 to 12, q: an integer of from 2 to 6). F(CF₂)_m-, (CF₃)₂CF(CF₂)_m-, H(CF₂)_m-, F(CF₂)_m(CH₂)_q-, (CF₃)₂CF(CF₂)_m(CF₂)_q-, (CF₃)₂C(CH₃)(CH₂)_q-, CF₃CHF(CF₂)(CH₂)_q-, H(CF₂)_m(CH₂)_q-, (CF₃)₂C=CF-, F(CF₂)_mCH=CH(CH₂)_q-, F(CF₂)_mCH₂CHI(CH₂)_q-, (CF₃)₂CF(CF₂)_mCH₂CHI(CH₂)_q-, F(CF₂)_mCH₂CH(OH)(CH₂)_q-, (CF₃)₂CF(CF₂)_mCH₂CH(OH)(CH₂)_q-, CF₃O[CF(CF₃)CF₂O]_mCF(CF₃)-, CF₃CHFO(CF₂)₃-, F(CF₂)₃OCF₂(CF₃)(CH₂)_q-, (CF₃)₂CH-, and F(CF₂)_m(CH₂)_qO(CH₂)_q-.

Further, as the polymerizable monomer (a), a (meth)acrylic acid amide such as F(CF₂)_mCH₂NHCOCR'=CH₂ wherein R' is a hydrogen atom or a methyl group, may preferably be mentioned, in addition to the above.

In the fluorine-containing copolymer (B), the proportion of the above polymerizable monomer (a) to the

total polymerizable monomers, is preferably at least 20 mol%, more preferably from 20 to 95 mol%, particularly preferably from 30 to 90 mol%. If the above proportion is less than 20 mol%, no adequate surface anti-fouling property may be obtained, whereby fouling is likely to be remarkable during use for a long period of time.

Further, as the polymerizable monomer (b), a compound of the formula 2 is preferred:



wherein R^2 is a hydrogen atom or a C_{1-4} alkyl group, each of c and d which are independent of each other, is an integer of from 0 to 6, Z is a single bond or a group selected from the group consisting of $-\text{CH}_2-$, $-\text{O}-$, $-\text{COO}-$, $-\text{CONH}-$, $-\text{NHCO}-$ and $-\text{OCO}-$, and L is a photo-curable functional group having no thermal polymerizability.

L is preferably a cinnamoyl group. For example, a polymerizable monomer (b) wherein L is a cinnamoyloxy group, is obtainable by reacting an unsaturated compound corresponding to the formula 2 wherein L is replaced by a hydroxyl group, with cinnamoyl chloride.

Specifically, the polymerizable monomer (b) may, for example, be a cinnamic acid ester of an allyl alcohol-alkylene oxide adduct, a cinnamic acid ester of 2-hydroxyethyl acrylate, or allyl cinnamate.

In the fluorine-containing copolymer (B), the proportion of the above polymerizable monomer (b) to the total polymerizable monomers, is preferably at least 5

mol%, more preferably from 5 to 80 mol%, still more preferably from 10 to 70 mol%, most preferably from 10 to 50 mol%. If the above proportion is less than 5 mol%, it tends to be difficult to form a photo-cured coating film having adequate hardness, and fouling is likely to be remarkable during use for a long period of time.

Further, the fluorine-containing copolymer (b) may be one obtained by copolymerizing another polymerizable monomer (c) in addition to the above-described polymerizable monomers (a) and (b).

The polymerizable monomer (c) is preferably a polymerizable monomer having no polyfluoroalkyl group or photo-curable functional group. Specifically, a polymerizable monomer having a polyoxyalkylene chain, such as polyoxypropylene glycol monoalkyl ether (meth)acrylate, having a molecular weight of from 100 to 3,000, may be mentioned.

Further, the following compounds may also be mentioned.

Methyl vinyl ether, ethyl vinyl ether, butyl vinyl ether, isobutyl vinyl ether, cyclohexyl vinyl ether, chloroethyl vinyl ether,

ethylene, propylene, 1-butene, isobutylene, cyclohexene, styrene, α -methylstyrene,

methyl allyl ether, ethyl allyl ether, butyl allyl ether, cyclohexyl allyl ether,

vinyl acetate, vinyl propionate, vinyl butyrate,

vinyl isobutyrate, vinyl valerate, vinyl caproate, vinyl
caprylate, Veova 9 and Veova 10 (tradenames, manufactured
by Shell Chemical) (vinyl esters of C₉ and C₁₀ branched
fatty acids), vinyl versatate, allyl acetate,

5 methyl (meth)acrylate, ethyl (meth)acrylate, butyl
(meth)acrylate, cyclohexyl (meth)acrylate,

ethyl crotonate, butyl crotonate, cyclohexyl
crotonate, (meth)acrylic acid amide, acrylonitrile, 2,4-
dicyanobutene-1, isoprene, butadiene,

10 vinyl chloride, vinylidene chloride,
trifluorochloroethylene, tetrafluoroethylene, vinylidene
fluoride, etc.

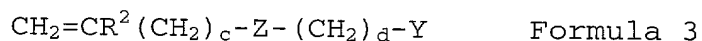
Among the above, preferred is a compound having a
C₁₋₁₅ linear, branched or alicyclic alkyl group.

15 By using the polymerizable monomer (c), there will
be effects for improving the storage stability of the
curable composition in addition to the effects for
improvement of the film-forming property of the fluorine-
containing copolymer (B), improvement of the
20 compatibility and dispersibility of the fluorine-
containing copolymer (B) to the above compound (A) and
improvement of e.g. a surface migration property of the
curable composition.

In the fluorine-containing copolymer (B), the
25 proportion of the above polymerizable monomer (c) to the
total polymerizable monomers, is preferably at most 75
mol%. Namely, it is preferred that the total proportion

of the above polymerizable monomers (a) and (b) to the
total polymerizable monomers, is at least 25 mol%. If
the total proportion is less than 25 mol%, no adequate
weather resistance or surface anti-fouling property can
5 be obtained.

As the polymerizable monomer (d), a compound of the
following formula 3 may, for example, be mentioned:



wherein R^2 is a hydrogen atom or a C_{1-4} alkyl group, each
10 of c and d which are independent of each other, is an
integer of from 0 to 6, Z is a single bond or a group
selected from the group consisting of $-\text{CH}_2-$, $-\text{O}-$, $-\text{COO}-$,
 $-\text{CONH}-$, $-\text{NHCO}-$ and $-\text{OCO}-$, and Y is a group having a group
capable of introducing a photo-curable functional group.

15 Y is preferably a C_{1-20} substituted or non-substituted
organic group having a hydroxyl group, an epoxy group, an
amino group, an isocyanate group or a carboxyl group,
particularly preferably an organic group having a
hydroxyl group or an epoxy group.

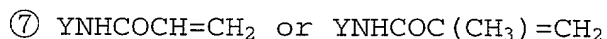
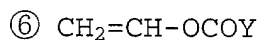
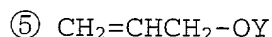
20 The following compounds may be mentioned as the
polymerizable monomer (d). In the following formulae, Y
has the same meaning as mentioned above.

① a polymerizable carboxylate such as a
(meth)acrylate

25 ② $\text{CH}_2=\text{CH}-\text{COOY}$

③ $\text{CH}_2=\text{C}(\text{CH}_3)-\text{COOY}$

④ $\text{CH}_2=\text{CH}-\text{OY}$



Specifically, the following compounds may preferably
5 be mentioned as the polymerizable monomer (d).

(Meth)acrylic acid, 2-hydroxyethyl (meth)acrylate,
3-hydroxypropyl (meth)acrylate, 3-phenoxy-2-hydroxypropyl
(meth)acrylate, 3-chloro-2-hydroxypropyl-(meth)acrylate,
(2-hydroxyethyl) allyl ether, N-methylolacryloamide,
10 mono(2-methacryloyloxyethyl) succinate, mono(2-
methacryloyloxyethyl) phthalate, glycidyl (meth)acrylate,
(meth)allylglycidyl ether, $\text{CH}_2=\text{CH}-\text{COOCH}_2\text{CH}_2\text{PO}(\text{OH})_2$,
 $\text{CH}_2=\text{C}(\text{CH}_3)-\text{COOCH}_2\text{CH}_2\text{PO}(\text{OH})_2$, etc.

In a case where the fluorine-containing copolymer
15 (B) is a compound obtainable by introducing a photo-
curable functional group into a fluorine-containing
copolymer (D) obtained by copolymerizing the above
polymerizable monomer (a) and the above polymerizable
monomer (d), the photo-curable functional group is
20 preferably a (meth)acryloyl group, a cinnamoyl group or
an azide group.

As a method of introducing a photo-curable
functional group into the above fluorine-containing
copolymer (D), for example, the fluorine-containing
25 copolymer (D) having a hydroxyl group, may be reacted
with acryl chloride or cinnamoyl chloride to introduce an
acryloyloxy group or a cinnamoyloxy group. Here, the

group capable of introducing the photo-curable functional group may remain in the fluorine-containing copolymer (B) obtained by introducing the photo-curable functional group into the fluorine-containing copolymer (D). In
5 such a case, the dispersibility of the fluorine-containing copolymer (B) to the above compound (A) will be improved, and the film-forming property of the fluorine-containing copolymer (B) will be improved.

The fluorine-containing copolymer (B) to be used in
10 the present invention, may, for example, be obtained by the following methods.

(1) The above polymerizable monomer (a) and the above polymerizable monomer (b) are copolymerized. If necessary, the above polymerizable monomer (c) may also
15 be copolymerized.

(2) The above polymerizable monomer (a) and the above polymerizable monomer (d) are copolymerized to obtain a fluorine-containing copolymer (D). Then, a photo-curable functional group is introduced into at
20 least a part of functional groups capable of introducing a photo-curable functional group, of the fluorine-containing copolymer (D). Further, at the time of obtaining the fluorine-containing copolymer (D), the above polymerizable monomer (c) may be copolymerized, as
25 the case requires.

The polymerization method is not particularly limited, and a method such as solution polymerization,

emulsion polymerization, suspension polymerization or bulk polymerization, may be employed. The polymerization can be carried out by applying a polymerization initiating source such as a polymerization initiator or an ionizing radiation to predetermined amounts of the above-mentioned various polymerizable monomers. Further, the molecular weight may be adjusted by adding a suitable chain transfer agent, or the polymerization can be carried out in the presence of the above compound (A).

Other various conditions may be the same as the conditions which are commonly employed in usual solution polymerization, emulsion polymerization, suspension polymerization or bulk polymerization.

The molecular weight of the fluorine-containing copolymer (B) is preferably from 300 to 100,000, more preferably from 1,000 to 50,000. Especially when the curable composition of the present invention is used without using any solvent (e.g. as a sealing material), if the molecular weight of the fluorine-containing copolymer (B) is large, the working efficiency tends to be very poor. Accordingly, the molecular weight is preferably at most 15,000, particularly preferably at most 10,000.

The amount of the fluorine-containing copolymer (B) in the curable composition of the present invention, is preferably from 0.01 to 20 parts by mass, particularly preferably from 0.3 to 7 parts by mass, per 100 parts by

mass of the compound (A).

Now, the photopolymerization initiator (C) will be described.

As the photopolymerization initiator, a known
5 initiator such as an arylketone type photopolymerization
initiator (such as an acetophenone, a benzophenone, an
alkylaminobenzophenone, a benzyl, a benzoine, a benzoine
ether, a benzyl dimethylketal, a benzoyl benzoate or an
 α -acyloxime ester), a sulfur-containing
10 photopolymerization initiator (such as a sulfide or a
thioxanthone), an acylphosphine oxide type
photopolymerization initiator or a diacylphosphine oxide
type photopolymerization initiator, may preferably be
employed. Further, a plurality of photopolymerization
15 initiators may be used in combination.

In a case where a cured coating film of the curable
composition of the present invention is to be used as a
protective film for a photomask, it is preferred to use a
photopolymerization initiator having an absorptivity (e)
20 per mol of at most 100 (ℓ /mol/cm) at 365 nm. The
absorptivity is a coefficient to show the degree of light
absorption by a substance and is a constant specific to
the substance. If the light intensity becomes from I_0 to
I when light passes through a substance layer having a
25 thickness of s, e represented by the following formula,
is the absorptivity (unit: ℓ /mol/cm).

$$I=I_0 \times 10^{(-es)}$$

Accordingly, in the above case, an acetophenone
type, benzoine type or benzophenone type
photopolymerization initiator is preferred from the
viewpoint of the transmittance of light with a wavelength
of 365 nm. Particularly preferred is an acetophenone
type photopolymerization initiator. Specifically,
compounds disclosed in paragraphs 0063 to 0065 in JP-A-
11-268196, may be mentioned.

The amount of the photopolymerization initiator (C)
in the curable composition of the present invention, is
preferably from 0.01 to 20 parts by mass, particularly
preferably from 0.1 to 10 parts by mass, per 100 parts by
mass of the compound (A).

The curable composition of the present invention may
further contain the following solvent and various
additives, in addition to the above-described basic
components.

Solvent

The curable composition of the present invention may
contain a solvent for the purpose of adjusting the
viscosity of the composition or improving the storage
stability. The solvent may, for example, be an aliphatic
hydrocarbon, an aromatic hydrocarbon, a halogenated
hydrocarbon, an alcohol, a ketone, an ester, an ether, an
ester alcohol, a ketone alcohol, an ether alcohol, a
ketone ether, a ketone ester or an ester ether.
Particularly, it is preferred to use an alcohol, whereby

the storage stability will be improved.

As the alcohol, a C₁₋₁₀ alkyl alcohol is preferred, and methanol, ethanol, isopropyl alcohol, isoamyl alcohol or hexyl alcohol may, for example, be mentioned.

5 The amount of the solvent is preferably at most 500 parts by mass, per 100 parts by mass of the total amount of the above compound (A) and the above fluorine-containing copolymer (B).

Adhesion-improving agent

10 The curable composition of the present invention may contain an adhesion-improving agent such as a silane coupling agent or polysilazane in order to improve the adhesion between the cured coating film and the substrate.

15 As the silane coupling agent, a known compound may be employed. In the present invention, a compound having a functional group such as a (meth)acryloyloxy group, an amino group, a mercapto group, an epoxy group or an isocyanate group, is preferred. Specifically, silanes
20 disclosed in paragraphs 0068 to 0072 in JP-A-11-240104, may be mentioned.

As a method of use, there may be mentioned a method wherein the adhesion-improving agent is added to the curable composition, or a method wherein the adhesion-
25 improving agent is preliminarily coated on the surface of a substrate to form a same layer, and then the curable composition is coated thereon.

Dehydrating agent

To the curable composition of the present invention, a dehydrating agent may be added to further improve the storage stability of the composition.

5 The dehydrating agent may, for example, be ① an alkyl orthoformate such as methyl orthoformate or ethyl orthoformate, ② an alkyl orthoacetate, such as methyl orthoacetate or ethyl orthoacetate, ③ a hydrolyzable organic silicon compound such as methyltrimethoxysilane, 10 vinyl trimethoxysilane, tetramethoxysilane or tetraethoxysilane, or ④ a hydrolyzable organic titanium compound. In the present invention, vinyl trimethoxysilane or tetraethoxysilane is particularly preferred from the viewpoint of the cost and effects.

15 The amount of the dehydrating agent is preferably at most 30 parts by mass, per 100 parts by mass of the total amount of the compound (A) and the fluorine-containing copolymer (B). If the amount of the dehydrating agent exceeds the above range, adverse effects to the hardness 20 and flexibility of the curable composition, are likely to be brought about.

Age resistor

As an age resistor, an antioxidant, an ultraviolet absorber or a photostabilizer, which is commonly 25 employed, may be used.

As the ultraviolet absorber, a known ultraviolet absorber can be used. For example, an ultraviolet

absorber of benzotriazole type, benzophenone type,
salicylic acid type or phenyltriazine type, may be
mentioned. Specifically, in addition to the compound
disclosed in paragraph 0078 in JP-A-11-268196, 2-hydroxy-
5 3-methacryloyloxypropyl-3-(3-(benzotriazole-2-yl)-4-
hydroxy-5-t-butylphenyl)propionate, 2-[4-(2-hydroxy-3-
dodecyloxypropyloxy)-2-hydroxyphenyl]-4,6-bis(2,4-
dimethylphenyl)-1,3,5-triazine, 2-[4-(2-hydroxy-3-
tridecyloxypropyloxy)-2-hydroxyphenyl]-4,6-bis(2,4-
10 dimethylphenyl)-1,3,5-triazine, and 2-(4-hexyloxy-2-
hydroxyphenyl)-4,6-diphenyl-1,3,5-triazine, may, for
example, be mentioned.

In the present invention, one having a polymerizable
functional group in its molecule, such as 2-{2-hydroxy-5-
15 (2-acryloyloxyethyl)phenyl}benzotriazole or 2-hydroxy-3-
methacryloyloxypropyl 3-(3-(benzotriazole-2-yl)-4-
hydroxy-5-t-butylphenyl) propionate, is particularly
preferred.

As the photostabilizer, a hindered amine type
20 photostabilizer which is used as a photostabilizer for a
synthetic resin, may preferably be mentioned.
Specifically, the compounds disclosed in paragraph 0080
in JP-A-11-268196 may be mentioned.

The antioxidant may, for example, be a hindered
25 phenol type antioxidant such as 2,6-di-t-butyl-p-cresol,
or a phosphorus type antioxidant such as triphenyl
phosphite.

Others

As other additives, a leveling agent such as a silicon resin type leveling agent or an acrylic resin type leveling agent, and a silicon resin type defoaming agent such as polydimethylsiloxane, may be used.

Now, the substrate will be described.

The substrate to be used in the present invention is not particularly limited, and various synthetic resins may be employed, such as an aromatic polycarbonate, a polymethyl methacrylate, a polystyrene, an aromatic polyester such as polyethylene terephthalate, a polyarylate, a polyimide, a polyethersulfone, a polysulfone, a polyamide, a cellulose triacetate, a fluororesin, an acrylic resin, a diethylene glycol bisallylcarbonate resin (such as CR-39), a copolymer of dimethacrylate having a halogen-substituted aromatic ring with a monofunctional monomer having an aromatic ring, a copolymer of a polyisocyanate with a polythiol, a methyl methacrylate/styrene resin, a tricyclo[5.2.1.0^{2,6}]deca-8-yl methacrylate/styrene resin, a copolymer of tricyclo[5.2.1.0^{2,6}]deca-8-yl methacrylate/styrene/a crosslinkable polyfunctional monomer, a copolymer of an aromatic ring-containing di(meth)acrylate/an aromatic ring-containing monomer/a hydroxyl group-containing monomer, a copolymer of an aromatic ring-containing di(meth)acrylate/an aromatic ring-containing monomer/an epoxy group-containing monomer, a copolymer of a

di(meth)acrylate containing a halogen-substituted
aromatic ring and an alkylene glycol group/an aromatic
ring-containing monomer/a compound containing an aromatic
ring and an epoxy group, and a copolymer of a styrene
5 derivative/ethylene glycol dimethacrylate/a specific
diacrylate compound/a specific epoxy-modified
di(meth)acrylate.

In the present invention, a photomask (an emulsion
mask) is preferred, which is prepared by using an
10 emulsion dry plate having a photosensitive emulsion
prepared by dispersing a photosensitive agent of silver
nitrate type or silver halide type coated on a substrate
made of e.g. glass or a transparent resin, such as a
polyester such as polyethylene terephthalate, an acryl
15 resin or a polycarbonate.

A coated substrate (such as a photomask) having a
cured coating film formed by using the curable
composition of the present invention, is obtainable as
follows.

20 By means of an exposure apparatus (such as a
photoplotter), a predetermined pattern is drawn and
recorded on an emulsion dry plate, and the drawn emulsion
dry plate is subjected to development treatment
(development, fixing, washing with water, drying),
25 followed by a correcting and inspecting step to obtain a
photomask original plate. On the surface of the emulsion
layer of the photomask original plate, the above-

mentioned curable composition is uniformly coated and
dried to remove the solvent contained in the curable
composition, whereupon an active energy ray is irradiated
to cure the curable composition, to form a cured coating
5 film (protective layer).

The method of coating the curable composition of the
present invention on the surface of the substrate, is not
particularly limited, and a die-coating method, a spin
coating method, a dip coating method, a flow coating
10 method, a roll coating method or a spray coating method
may, for example, be employed. In the present invention,
a spray coating method or a die-coating method is
preferably employed from the viewpoint of the
productivity and the surface appearance.

15 Further, the active energy ray to cure the curable
composition of the present invention is not particularly
limited, and an ultraviolet ray, an electron ray or other
active energy rays may be employed. However, an
ultraviolet ray is particularly preferred, since the
20 apparatus is simple, and the productivity is excellent.
As an ultraviolet ray source, a xenon lamp, a pulse xenon
lamp, a low pressure mercury lamp, a high pressure
mercury lamp, a super high pressure mercury lamp, a metal
halide lamp, a carbon arc lamp or a tungsten lamp, may,
25 for example, be used.

The thickness of the cured coating film formed by
the curable composition of the present invention is

preferably from 0.1 to 50 μm , more preferably from 0.3 to 30 μm . If the thickness of the cured coating film exceeds 50 μm , curing by an active energy ray tends to be inadequate, whereby the adhesion with the substrate tends to be poor. On the other hand, if it is less than 0.1 μm , the abrasion resistance and scratch resistance of the cured coating film tends to be inadequate.

Now, the present invention will be described in detail with reference to Examples. However, it should be understood that the present invention is by no means restricted thereto.

As the substrate, a transparent aromatic polycarbonate resin plate (thickness: 3 mm) was used in Examples 1 to 8, and a photomask original plate prepared by drawing a predetermined pattern on a usual emulsion dry plate by a usual method, followed by development treatment, was used in Examples 9 and 10. Further, measurements and evaluations of various properties with respect to the samples obtained in the respective Examples were carried out by the following methods, and the results are shown in Tables 1 and 2.

Abrasion resistance

In accordance with the abrasion resistance test method of JIS R3212, two CS-10F truck wheels were combined with a weight of 500 g, respectively, and rotated 100 times, whereby the haze was measured by a haze meter, and the abrasion resistance was represented

by the change in the haze between before and after the abrasion test.

Scratching resistance (pencil hardness)

In accordance with JIS K5400, the test was carried
5 out by means of a pencil scratching tester.

Adhesion properties

The test was carried out in accordance with the cross cut test of JIS K5400. The number (m) of cross cut sections where the coated film remained without being
10 peeled, is represented by m/100.

Contact angle (surface tension)

The contact angle of a water droplet of distilled water dropped on the substrate surface to have a diameter of 1 mm, was measured. Measurements were carried out at
15 five different places on the substrate surface, and the average value was calculated.

Anti-fouling property (finger print-wiping efficiency)

A finger was pressed on the substrate surface to leave a finger print, and then, the finger print was
20 wiped off with a tissue paper, and the degree of removal of the sebum was visually judged.

Non-tackiness (adhesive tape peeling)

A commercially available adhesive tape was closely bonded to the substrate surface, whereupon the peeling
25 property was qualitatively judged.

Weather resistance

By means of a sunshine weatherometer, a sample was

subjected to exposure for 500 hours with a cycle of rain falling for 12 minutes and drying for 48 minutes at a black panel temperature of 63°C, whereupon the contact angle was measured.

5 PREPARATION EXAMPLE 1

Using n-butanol as an initiator, polymerization of propylene oxide was carried out in the presence of a zinc hexacyanocobaltate catalyst to obtain a polypropylene oxide monobutyl ether (average molecular weight: 2,000).

10 300 g (0.15 mol) of this compound was put into a 2 l flask, and further, 500 g of acetone, 11.7 g (0.17 mol) of triethylamine and 150 mg of hydroquinone monomethyl ether were added, followed by stirring at 40°C for 10 minutes in a nitrogen stream.

15 Then, a solution having 14.5 g (0.16 mol) of acrylic acid chloride dissolved in 100 g of acetone, was dropwise added over a period of 30 minutes. Then, the temperature was raised to 57°C, followed by stirring for one hour and thirty minutes to complete the reaction. Then, the
20 ammonium salt was filtered off, and the filtrate was dropwise added to 3 l of water with stirring to reprecipitate the compound, whereupon the precipitated compound was washed twice with water and then dried to obtain a polymerizable monomer (c1) having an acryloyl
25 group and a polyether chain.

PREPARATION EXAMPLE 2

In the same manner as in Preparation Example 1, a

polypropylene oxide monobutyl ether (average molecular weight: 400) was obtained. And, acrylic acid chloride was added thereto to change the terminal hydroxyl group to an acryloyl group, to obtain a polymerizable monomer (c2) having an acryloyl group and a polyether chain.

PREPARATION EXAMPLE 3

Into a 500 ml flask, 8.7 g (0.15 mol) of allyl alcohol was put, and further, 100 g of acetone, 17.7 g (0.17 mol) of triethylamine and 15 mg of hydroquinone monomethyl ether were added, followed by stirring at 40°C for 10 minutes in a nitrogen stream. Then, a solution having 26.6 g (0.16 mol) of cinnamoyl chloride dissolved in 100 g of acetone, was dropwise added over a period of 30 minutes. Then, the temperature was raised to 57°C, followed by stirring for one hour and thirty minutes to complete the reaction. Then, the ammonium salt was filtered off, and the filtrate was dropwise added to 3 l of water with stirring to reprecipitate the compound, whereupon the precipitated compound was washed twice with water and then dried to obtain a polymerizable monomer (b1) having a cinnamoyl group and an allyl group.

PREPARATION EXAMPLE 4

Into a 500 ml flask, 46.16 g of the above polymerizable monomer (c1), 23.91 g of $C_tF_{2t+1}CH_2CH_2OCOCH=CH_2$ (average value of t: 9) (hereinafter referred to as a polymerizable monomer (a1)) and 5.36 g of 2-hydroxyethyl acrylate, were dissolved in a solvent

10000019 "120404"
mixture comprising 150 g of chloroform and 150 g of
isopropyl alcohol, and 1.5 g of azoisobutyronitrile was
added as an initiator, followed by thermal polymerization
at 70°C for 10 hours. Thereafter, the solvent was
5 removed to obtain a fluorine-containing polymer (d1)
having no photo-curable functional group. Then, in the
same manner as in Preparation Example 2, an acryloyl
group was introduced into the fluorine-containing polymer
(d1) to obtain a fluorine-containing copolymer (b1)
10 (acryloyl group content: 20 mol%, mass average molecular
weight: 25,000).

EXAMPLE 1

Into a 200 ml four-necked flask equipped with a
stirrer and a condenser, 60 g of butyl acetate, 20 g of
15 isopropyl alcohol, 10 g of dipentaerythritol
hexaacrylate, 10 g of tris(acryloxyethyl) isocyanurate,
0.6 g of 2-methyl-1-{4-(methylthio)phenyl}-2-
morpholinopropane-1-one, 0.5 g of 2-{2-hydroxy-5-(2-
acryloyloxyethyl)phenyl}benzotriazole and 0.1 g of a
20 silicone type leveling agent "BYK306" (tradename,
manufactured by Bic Chemy Company, the same applies
hereinafter), were added and dissolved, and further, 0.2
g of N-(2-aminoethyl)-3-aminopropyltrimethoxysilane and
0.2 g of the fluorine-containing copolymer (B1) were
25 added, followed by stirring at room temperature for 1
hour to obtain a coating solution 1.

Then, the coating solution 1 was coated on a

substrate by a die-coating method (wet thickness: 10 μm)
and dried for 5 minutes in an oven of 90°C, and then
irradiated with an ultraviolet ray by means of a high
pressure mercury lamp with 500 mJ/cm^2 (accumulated energy
amount of ultraviolet ray having a wavelength within a
range of from 300 to 390 nm, the same applies
hereinafter) in an air atmosphere to obtain a sample
having a cured coating film formed in a thickness of 2
 μm .

10 EXAMPLES 2 to 6

Respective samples were obtained in the same manner
as in Example 1 except that the following fluorine-
containing copolymers (B2) to (B6) were used,
respectively, instead of the fluorine-containing
15 copolymer (B1).

Example 2: A coating solution 2 was prepared by
using a fluorine-containing copolymer (B2) (acryloyl
group content: 27 mol%, mass average molecular weight:
30,000) obtained by copolymerizing the above
20 polymerizable monomer (a1), polymerizable monomer (c2)
and 2-hydroxyethyl acrylate in a molar ratio of 40/30/30
and then changing the hydroxyl group in the molecule to
an acryloyl group by means of acrylic acid chloride.

Example 3: A coating solution 3 was prepared by
25 using a fluorine-containing copolymer (B3) (acryloyl
group content: 36 mol%, mass average molecular weight:
9,000) obtained by copolymerizing the above polymerizable

monomer (a1), stearyl acrylate and 2-hydroxyethyl
acrylate in a molar ratio of 35/25/40 and then changing
the hydroxyl group in the molecule to an acryloyl group
by means of acrylic acid chloride.

5 Example 4: A coating solution 4 was prepared by
using a fluorine-containing copolymer (B4) (acryloyl
group content: 14 mol%, mass average molecular weight:
9,000) obtained by copolymerizing the above polymerizable
monomer (a1), butyl acrylate and 2-hydroxyethyl acrylate
10 in a molar ratio of 35/50/15 and then changing the
hydroxyl group in the molecule to an acryloyl group by
means of acrylic acid chloride.

Example 5: A coating solution 5 was prepared by
using a fluorine-containing copolymer (B5) (acryloyl
15 group content: 20 mol%, mass average molecular weight:
12,000) obtained by copolymerizing the above
polymerizable monomer (a1), polymerizable monomer (b1)
and stearyl acrylate in a molar ratio of 40/20/40.

Example 6: A coating solution 6 was prepared by
20 using a fluorine-containing copolymer (B6) (acryloyl
group content: 20 mol%, mass average molecular weight:
8,000) obtained by copolymerizing the above polymerizable
monomer (a1), polymerizable monomer (b1) and butyl
acrylate in a molar ratio of 40/20/40.

25 EXAMPLE 7

A coating solution 7 was prepared in the same manner
as in Example 1 except that no fluorine-containing

copolymer (B1) was contained, and a sample was prepared
by using this coating solution.

EXAMPLE 8

A coating solution 8 was prepared in the same manner
5 as in Example 1 except that the fluorine-containing
copolymer (D1) was used instead of the fluorine-
containing copolymer (B1), and a sample was obtained by
using this coating solution.

100004-100004

Table 1

	Abrasion resistance	Adhesion	Contact angle	Anti- fouling property	Non- tackiness	Weather resistance	Appearance
Example 1	5.5	100/100	108°	1 to 2 times	Good	102°	Good
Example 2	6.2	100/100	108°	1 to 2 times	Good	100°	Good
Example 3	5.7	100/100	108°	1 to 2 times	Good	98°	Good
Example 4	4.8	100/100	108°	1 to 2 times	Good	105°	Good
Example 5	6.5	100/100	108°	1 to 2 times	Good	101°	Good
Example 6	5.9	100/100	108°	1 to 2 times	Good	100°	Good
Example 7	4.9	100/100	73°	5 to 6 times	Poor	65°	Good
Example 8	7.8	100/100	108°	2 to 3 times	Good	75°	Good

EXAMPLE 9

Into a 200 ml four-necked flask equipped with a stirrer and a condenser, 80 g of 2-butanone, 10 g of dipentaerythritol hexaacrylate, 10 g of tris(acryloxyethyl) isocyanurate, 0.6 g of 2-methyl-1-(4-(methylthio)phenyl)-2-morpholinopropan-1-one, and 0.1 g of a silicone type leveling agent "BYK306", were added and dissolved, and further, 0.2 g of N-(2-aminoethyl)-3-aminopropyltrimethoxysilane and 0.2 g of the fluorine-containing copolymer (B1) were added, followed by stirring at room temperature for 1 hour to obtain a coating solution 9.

Then, the coating solution 9 was coated on a substrate by a die-coating method (wet thickness: 10 μ m) and dried for 5 minutes in an oven of 90°C, and then irradiated with a ultraviolet ray by means of a high pressure mercury lamp with 500 mJ/cm² in an air atmosphere, to obtain a sample having a cured coating film formed in a thickness of 2 μ m.

EXAMPLE 10

The substrate having no cured coating film formed, was used as it is, as a sample.

Table 2

	Pencil hardness	Adhesion	Contact angle	Non-tackiness	Appearance
Example 1	3H	100/100	108°	Good	Good
Example 2	4B	-	65°	Poor	Good

According to the present invention, it is possible to provide a curable composition capable of forming in a single coating step, a cured coating film which has an anti-fouling property and a release property and is excellent in abrasion resistance, whereby such properties will not deteriorate even when used for a long period of time, a cured coating film having the above-mentioned properties, formed by means of such a curable composition, and a coated substrate having such a cured coating film. The cured coating film is most suitable as a protective layer for a photomask.

The entire disclosures of Japanese Patent Application No. 2000-375748 filed on December 11, 2000 and Japanese Patent Application No. 2001-327644 filed on October 25, 2001 including specifications, claims and summaries are incorporated herein by reference in their entireties.